STEWART LAKE WATERFOWL MANAGEMENT AREA

REPORT ON THE SEDIMENT SAMPLES COLLECTED FROM STEWART LAKE, UTAH DURING 2002

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JANUARY 2004

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Report Summary

This report addresses sediment samples collected by personnel of the Bureau of Reclamation in April, August, September, and October 2002 at Stewart Lake, Utah, in conjunction with the effort to remediate its selenium contaminated sediments. The April samples were collected as a pre-flood baseline. Surface samples were collected for analysis of total selenium from 20 permanent monitoring sites. Sediment depth profile samples were collected from 4 quadrants in site 21 for selenium speciation; surface samples were collected at 5 other sites for speciation. Because of low runoff in the spring of 2002, no flood water entered Stewart Lake from the Green River. Samples collected in August and September related to a 5-day (August 29 through September 2) controlled flood that was confined to 2 experimental plots constructed in the north end of Stewart Lake. Samples were collected for total selenium and speciation. The October samples consisted of the routine surface samples collected for total selenium analysis annually at 20 permanent monitoring sites in the lake.

At the 5 sites sampled for selenium speciation at the surface only, selenate was present in its highest at site 15, 25 percent of the total; it was between 10 and 20 percent at sites 7 and 10; and it was lowest at sites 1 and 16 at less than 5 percent. Site 16 had been covered by a pool of ice formed by seepage shortly before sampling. Elemental selenium was predominant at 3 of the 5 sites, while selenite dominated at the other 2 sites. Elemental selenium was the dominant species in all surface samples from site 21, where it constituted between 40 and 50 percent of the total selenium. However, selenate was approximately 25 percent of the total in 3 of the 4 quadrants before lime was applied.

Lime was to be applied to the test plot at site 21 before the spring flood from the Green River in 2002. Samples were collected before and after the lime was applied to the 3-acre experimental plot. Application of the lime caused chemical reduction of oxidized species of selenium (selenate and selenite) and a 1-2 ppm increase in reduced (elemental) selenium. It also caused a reduction in total selenium in 3 of the 4 sites sampled; the reductions amounted to between 1.6 and 3 ppm. The lime application also resulted in a net reduction in soluble selenium, indicating that the application of lime would not result in additional leaching of selenium as had been planned.

The flood experiment was generally successful. There was a reduction in selenate in all layers of the sediments, although there was an indication that some had been converted to selenite and elemental selenium. Total selenium was reduced in the surface sediments and in layers between 12 and 24 inches deep, but increased in the 6-12 inch deep layer. There was no large increase in selenium in deeper sediments that would indicate that there had been selenium deposition.

A separate analysis of the difference in the results of the flooding experiment based on total selenium as analyzed in laboratories at the University of California at Riverside and the USGS in Denver indicated that the analytical results differ. The USGS analytical procedure is based on a more vigorous digestion. The USGS procedure gives progressively higher results as selenium concentrations increase. In other words, the difference in the 2 sets of results is concentration dependent.

The change in the mass of selenium in the 2 experimental flood plots was estimated. In both the small experimental flood plot and 1 quadrant of the larger experimental flood plot (site 21), there was a net removal of selenium. There was a removal of soluble selenium in all layers (6-inch layers to a depth of 36 inches) of the sediment at each site. There was also a significant decrease in insoluble selenium in 4 of the 6 layers at site 21 and a net decrease in the profile as a whole. There were layers of apparent deposition of insoluble selenium at each site. The layers of deposition appear to be primarily due to the timing of the end of the flooding during the experiment. Estimates of the mass change in the surface sediments (0-6 inches) of the small flood plot and 4 quadrants of the large flood plot that were sampled separately showed a high degree of variability. In the surface sediments of the north half of site 21, there was a small net increase in selenium in 1 quadrant and a large decrease in another quadrant at site 21; there were relatively consistent decreases in the small flood plot and the 2 quadrants in the south half of site 21. An analysis of the quality control data from a 2-year period (2001-2002) indicated that much of the variability could be due to random variation.

The surface samples from the permanent monitoring sites indicate that there was a small increase (0.3 ppm average) in total selenium overwinter from November 2001 to April 2002. There was subsequently a small decrease (0.2 ppm average) between April and October 2002. Overall there was no change in total selenium during 2002.

The long-term monitoring data (1995-2002) were analyzed for trends. There have been overall decreases in total selenium in 1997 and 2000 and an overall increase during 1998. Since 1995, there has been no measurable overall change in total selenium at the permanent monitoring sites in Stewart Lake. Beginning in 1998, concentrations of total selenium have been much higher in the north end of Stewart Lake than to the south. Individual observations can be classified into appropriate locations within Stewart Lake with a near 90 percent accuracy on the basis of total selenium concentrations. Areas in the north of the lake have a total selenium concentration in excess of 20 ppm (average 28 ppm), while areas to the south in the lake have a selenium concentration that is less than 20 ppm (average 12 ppm). The increase in selenium that occurred in Stewart Lake in 1998 was confined to its north end. As was noted in the 2002 sediment report, the initial increase in 1998 was apparently caused by the J2, J3, and J4 drains before they were diverted to the Green River. The seepage has maintained the elevated levels since then. However, there has also been no change in total selenium at sites located in the south end of Stewart Lake away from the seepage. The total selenium concentration in these sites with lower selenium remains well above the selenium cleanup goal of 4 ppm.

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